AMENDMENT

In the Specification:

Prior to page 1, where the text of the application begins, please delete the double-sided cover page from the PCT stage if necessary.

At page 1, after the title, in the current single-paragraph that constitutes the entire Section of the Application pertaining to "Cross-reference to Related Applications", please delete the existing section and replace such deleted section with the following, two-paragraph replacement section:

The present application is a nationalization of International Patent Application PCT/AU99/00813, filed September 24, 1999, which claims priority to Australian Patent Application PP 6164, filed September 25, 1998.

FIELD OF THE INVENTION

This invention relates to methods for preparing cyclic peptides and peptidomimetics in solution and bound to solid supports, and to cyclic peptide or peptidomimetic libraries for use in drug screening programmes. In particular, the invention relates to a generic strategy for synthesis of cyclic peptides or peptidomimetics that enables the efficient synthesis under mild conditions of a wide variety of desired compounds.

After page 143, please start another page (144), and insert the following text of the Abstract, as taken from the priority application as filed and on first cover page from the PCT application:

This invention relates to methods for preparing cyclic peptides and peptidomimetic compounds in solution and bound to solid supports, and to cyclic peptide or peptidomimetic libraries for use in drug screening programmes. In particular, the invention relates to a generic

strategy for synthesis of cyclic peptides or peptidomimetics that enables the efficient synthesis under mild conditions of a wide variety of desired compounds. Two approaches were evaluated for their improvements in solution and solid phase synthesis of small cyclic peptides: positioning reversible *N*-amide substituents in the sequence; and applying native ligation chemistry in an intramolecular sense. Systematic investigation of the effects of preorganising peptides prior to cyclisation by using peptide cyclisation auxiliaries, and developing new linkers and peptide cyclisation auxiliaries to aid cyclic peptide synthesis gives surprising improvements in both yields and purity of products compared to the prior art methods. The combination of these technologies provides a powerful generic approach for the solution and solid phase synthesis of small cyclic peptides. The ring contraction and *N*-amide substitution technology of the invention provide improved methods for the synthesis of cyclic peptides and peptidomimetics. When used in conjunction with linker strategies, this combination provides solid-phase avenues to cyclic peptides and peptidomimetics.

At the appropriate pages, prior to the text on each page, please delete the header that reads "WO 00/18790 PCT/AU99/00813" if necessary.

In the Claims:

After entry into the U.S. national stage, and assurance of a U.S. filing date, please revise the claims from the enclosed PCT application as follows.

Please amend pending claims 4, 5, 6, 7, 8, 11, 13, 15, 16, 18, 19, 21, 22, 23, 24, 25, 26, 30, 31 and 36, so that the rewritten claims read as follows:

4. (Amended) A method according to claim 1, in which the cycle is a monocycle.